Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claim 1 (original): A hybrid polypeptide immunogen comprising a modified ORF0657n sequence segment at least about 100 amino acids in length, wherein said modified sequence segment comprises one or more alterations that increases sequence similarity to SEQ ID NO: 1.

Claim 2 (original): The hybrid polypeptide of claim 1, wherein said modified sequence segment comprises at least about 100 amino acids of a modified amino acid sequence selected from the group consisting of SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, and SEQ ID NO: 6, provided that said modified amino acid sequence contains at least 8 amino acid alterations that increase sequence similarity to SEQ ID NO: 1.

Claim 3 (original): The hybrid polypeptide of claim 3, wherein said modified amino acid sequence is SEQ ID NO: 2 containing 8 to 100 amino acid alterations that increase sequence similarity to SEQ ID NO: 1.

Claim 4 (currently amended): The hybrid polypeptide of claim 2, wherein said modified amino acid sequence has the following sequence:

X1-AIKNPAI-X2- DK-X3-H-X4-APN-X5- RPIDFEMK-X6-X7-X8-G-X9-QQFYHYAS-X10-V-X11- PARVIFT-X12-X13-K-X14-IELGLQ-X15-X16-X17-X18-W-X19-KFEVYEGDKKLP-X20- KLVSYD-X21-X22-KDYAYIRFSVSNGT-X23-X24-VKIVSSTH-X25-X26-X27-N-X28-X29-EKYDYTLM-X30- FAQPIYN-X31-X32-DK-X33-X34-X35- EEDY-X36-X37-X38- KLLAPYKKAKTLERQVY EL-X39- K-X40- Q-X41-KLPEKLKAEYKKKL-X42-X43-T-X44- KAL-X45-X46-QVKSA-X47- TEFQNV-X48-PTN-X49-K-X50- TDLQ-X51-X52-X53-X54-VV-X55-ESVEN-X56-ES-X57-MDTFV-X58-HPIKT-X59-X60-LNGKKY-X61-VM-X62- TTND-X63-YWKDF-X64- VEG-X65- RVRT-X66- SKD-X67- KNN-X68- RT-X69- IFPY-X70- EGK-X71-X72-YDAIVKV-X73- VKTI-X74-Y-X75-GOYHVRI-X76- DK-X77-X78-X79 (SEQ ID NO: 58)

wherein

X1 is either E or a D alteration;

X² is either K or an I alteration;

X³ is either D or an E alteration;

X⁴ is either S or a T alteration;

X⁵ is either S or a W alteration;

X6 is either K or an N alteration;

X7 is either K or a D alteration;

X8 is either D or a K alteration;

X9 is either T or an E alteration;

X10 is either S or a T alteration;

X¹¹ is either K or an E alteration;

X¹² is either D or a K alteration;

X13 is either S or a T alteration;

X14 is either E or an I alteration;

X15 is either S or a T alteration;

X16 is either G or an A alteration;

X17 is either K or a S alteration;

X18 is either F or a T alteration;

X19 is either R or a K alteration;

X²⁰ is either I or a V alteration;

X²¹ is either T or an S alteration;

X²² is either V or a D alteration;

X²³ is either K or an R alteration;

X²⁴ is either A or an E alteration;

X25 is either F or a Y alteration;

X26 is either an optionally present G insertion alteration;

X²⁷ is either N or a E alteration;

X²⁸ is either K or a I alteration

X²⁹ is either E or a H alteration;

X³⁰ is either E or a V alteration;

X³¹ is either S or a N alteration;

X32 is either A or a P alteration;

X33 is either F or an Y alteration;

X34 is either K or a V alteration;

X³⁵ is either T or a D alteration;

X³⁶ is either K or a N alteration;

X37 is either A or an L alteration;

X³⁸ is either E or a Q alteration;

X³⁹ is either N or an E alteration;

X⁴⁰ is either I or a L alteration;

X41 is either D or an E alteration;

X42 is either E or a D alteration;

X43 is either D or a Q alteration;

X44 is either K or an R alteration;

X45 is either D or an A alteration;

X⁴⁶ is either E or a D alteration;

X⁴⁷ is either I or a V alteration;

X⁴⁸ is either Q or a T alteration;

X⁴⁹ is either E or a D alteration;

X⁵⁰ is either M or an L alteration;

X51 is either D or an E alteration;

X52 is either T or an A alteration

X53 is either K or H alteration;

X⁵⁴ is either Y or an F alteration;

X55 is either Y or an F alteration;

X56 is either N or a S alteration;

X⁵⁷ is either M or a V alteration;

X58 is either K or an E alteration;

X⁵⁹ is either G or an A alteration;

X60 is either M or a T alteration;

X61 is either M or a V alteration;

X62 is either E or a K alteration;

X63 is either D or a S alteration;

X64 is either M or an I alteration;

X65 is either Q or a K alteration;

X66 is either I or a V alteration;

X67 is either A or a P alteration;

X68 is either T or an S alteration;

X69 is either I or a L alteration;

X⁷⁰ is either V or an I alteration;

X⁷¹ is either T or an A alteration;

X⁷² is either L or a V alteration;

X⁷³ is either H or a V alteration;

X⁷⁴ is either D or a G alteration;

X⁷⁵ is either D or an E alteration;

X⁷⁶ is either V or an I alteration;

X⁷⁷ is either E or a D alteration;

X⁷⁸ is either A or an I alteration;

X⁷⁹ is either F or a N alteration;

X¹ is either E or a D alteration;

X² is either K or an I alteration;

X³ is either D or an E alteration;

X⁴ is either S or a T alteration;

X⁵ is either S or a W alteration;

X6-X7-X8 is either KKD or NDK alterations;

X⁹-is-either T or an E alteration;

X¹⁰ is either S or a T alteration;

X¹¹ is either K or an E alteration;

X¹² is either D or a K alteration;

X¹³ is either S or a T alteration;

X14 is either E or an I alteration;

X15 is either S or a T alteration;

X¹⁶ is either G or an A alteration;

X¹⁷-X¹⁸ is either KF or ST alterations;

X¹⁹ is either R or a K alteration;

X²⁰ is either I or a V alteration;

X²¹ is either T or an S alteration;

X²² is either V or a D alteration;

X²³ is either K or an R alteration;

X²⁴ is either A or an E alteration;

X²⁵ is either F or a Y alteration;

X26-X²⁷ is either N or GE alterations;

X²⁸-X²⁹ is either KE or IH alterations;

X³⁰ is either E or a V alteration;

X³¹-X³² is either SA or NP alterations;

X³³ is either F or an Y-alteration; X³⁴-X³⁵ is either KT or VD alterations: X³⁶-X³⁷-X³⁸ is either KAE or NLO alterations: X³⁹ is either N or an E alteration: X⁴⁰ is either I or a L alteration; X⁴¹ is either D or an E alteration: X⁴² is either E or a D alteration; X⁴³ is either D or a O alteration: X⁴⁴ is either K or an R alteration: X⁴⁵-is either D or an A alteration; X46 is either E or a D alteration: X⁴⁷ is either I or a V alteration: X⁴⁸ is either Q or a T alteration; X⁴⁹ is either F or a D alteration: X⁵⁰ is either M or an L alteration: X⁵¹ is either D or an E alteration: X⁵²-X⁵³ is either TK or AH alterations: X⁵⁴ is either Y or an F alteration: X⁵⁵ is either Y or an F alteration; X⁵⁶ is either N or a S alteration: X⁵⁷ is either M or a V alteration: X⁵⁸ is either K or an E alteration: X⁵⁹ is either G or an A alteration; X⁶⁰ is either M or a T alteration: X61 is either M or a V alteration: X⁶² is either E or a K alteration: X⁶³ is either D or a S alteration; X64 is either M or an I alteration: X⁶⁵ is either O or a K alteration; X⁶⁶ is either I or a V alteration; X67 is either A or a P alteration: X⁶⁸ is either T or an S alteration; X⁶⁹ is either I or a L alteration: X⁷⁰ is either V or an Lalteration: X⁷¹-is either T or an A alteration:

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X<sup>72</sup> is either L or a V alteration;
X<sup>73</sup> is either H or a V alteration;
X<sup>74</sup> is either D or a G alteration;
X<sup>75</sup> is either D or an E alteration;
X<sup>76</sup> is either V or an I alteration;
X<sup>77</sup> is either E or a D alteration;
X<sup>78</sup> is either A or an I alteration;
X<sup>79</sup> is either F or a N alteration;
provided that at least 20 of said alterations are present.
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Claim 5 (original): The hybrid polypeptide of claim 4, wherein said modified sequence segment comprises at least 200 amino acids of said modified amino acid sequence.

Claim 6 (previously presented): The hybrid polypeptide of claim 4, wherein said modified sequence segment comprises said modified amino acid sequence and at least 55 of said alterations are present.

Claim 7 (original): The hybrid polypeptide of claim 1, wherein said hybrid polypeptide consists of a sequence selected from the group consisting of SEQ ID NOs: 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, and 43.

Claim 8 (original): A method of making a hybrid polypeptide comprising the step of introducing one or more alterations into a ORF0657n sequence segment at least about 100 amino acids in length, wherein at least one of said alterations increases sequence similarity to SEQ ID NO: 1.

Claim 9 (original): An immunogen comprising the modified ORF0657n sequence of claim 1 and one or more additional regions or moieties covalently joined to said sequence at the carboxyl terminus or amino terminus, wherein each region or moiety is independently selected from a region or moiety having at least one of the following properties: enhances the immune response, facilitates purification, or facilitates polypeptide stability.

Claim 10 (previously presented): A composition able to induce a protective immune response in a patient comprising an immunologically effective amount of the immunogen of claim 1 and a pharmaceutically acceptable carrier.

Claim 11 (original): The composition of claim 10, wherein said composition further comprises an adjuvant.

Claim 12 (previously presented): A method of inducing a protective immune response in a patient comprising the step of administering to said patient an immunologically effective amount of the immunogen of claim 1.

Claim 13 (original): The method of claim 12, wherein said patient is a human.

Claim 14 (original): The method of claim 13, wherein said patient is being treated prophylactically against *S. aureus* infection.

Claim 15 (previously presented): A nucleic acid comprising a nucleotide sequence encoding the polypeptide of claim 1.

Claim 16 (original): The nucleic acid of claim 15, wherein said nucleic acid is an expression vector and said nucleotide sequence is part of a recombinant gene.

Claim 17 (original): A cell comprising the recombinant gene of claim 16, wherein said recombinant gene expresses said nucleic acid sequence in said cell to produce said polypeptide.

Claim 18 (original): A method for evaluating the efficacy of an immunogen to produce a protective immune response against *Staphylococcus* comprising the steps of:

- (a) inoculating an animal model with said immunogen to produce an immunized animal model;
- (b) challenging said immunized animal model with a *Staphylococcus* challenge at a potency that provides about 80 to 90% death in said animal model over a period of about 7 to 10 days starting on the first or second day, wherein said *Staphylococcus* challenge is produced from *Staphylococcus* grown to stationary phase, and said *Staphylococcus* challenge is intravenously introduced into said immunized animal model; and

(c) measuring the ability of said immunogen to provide protective immunity.

Claim 19 (original): The method of claim 18, wherein said *Staphylococcus* is *Staphylococcus aureus*.

Claim 20 (original): The method of claim 19, wherein said animal model is a rat or mouse.

Claims 21-23 (Canceled).